

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification ⁶ : C02F 11/00</p>	<p>A1</p>	<p>(11) International Publication Number: WO 98/41479 (43) International Publication Date: 24 September 1998 (24.09.98)</p>
<p>(21) International Application Number: PCT/SE98/00444 (22) International Filing Date: 12 March 1998 (12.03.98) (30) Priority Data: 9700917-9 14 March 1997 (14.03.97) SE (71) Applicant (for all designated States except US): KEMIRA KEMI AB [SE/SE]; P.O. Box 902, S-251 09 Helsingborg (SE). (72) Inventors; and (75) Inventors/Applicants (for US only): HANSEN, Bengt [SE/SE]; Fridasgata 126, S-256 54 (SE). JOKINEN, Simo [FI/FI]; Sorsatie 19 B, FIN-90150 Oulu (FI). (74) Agent: AWAPATENT AB; P.O. Box 5117, S-200 71 Malmö (SE).</p>		<p>(81) Designated States: AL, AM, AT, AT (Utility model), AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, CZ (Utility model), DE, DE (Utility model), DK, DK (Utility model), EE, EE (Utility model), ES, FI, FI (Utility model), GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK (Utility model), SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).</p> <p>Published With international search report.</p>
<p>(54) Title: A METHOD FOR TREATING SLUDGE FROM WASTEWATER TREATMENT</p> <p>(57) Abstract</p> <p>A method for treating sludge from wastewater purification is described. In the method, a sludge is treated, which contains phosphorus and at least one metal which originates from precipitation chemicals and which is selected among divalent iron and aluminium, the pH of the sludge being adjusted to below 4, preferably below 2, for dissolving the content of phosphorus and said metal in the sludge; the remaining sludge is separated; the solution relieved of sludge and containing phosphorus and said metal is treated for precipitation of the phosphorus content of the solution as FePO_4 at a pH of 2-3; and precipitated FePO_4 is separated. The method is characterised in that the solution which remains after separation of FePO_4 and which contains said metal from the precipitation chemicals, is recycled to the wastewater purification. The phosphorus content of the solution is precipitated as FePO_4 by adding an at least equivalent amount of Fe^{3+}.</p> <div data-bbox="1088 1113 1429 1869"> <pre> graph TD SLUDGE --> I Hplus[H+] --> I I --> II II --> SLUDGE2[SLUDGE] II --> III S2[S2-] --> III III --> HMS[HMS] III --> IV OHminus[OH-] --> IV IV --> V Fe3plus[Fe3+] --> V V --> FePO4[FePO4] V --> Fe2Al3[Fe2+/Al3+] </pre> </div>		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakhstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

A METHOD FOR TREATING SLUDGE FROM WASTEWATER TREATMENT

The present invention relates to a method for treating sludge from wastewater treatment according to the preamble to claim 1. More specifically, the invention relates to such a process, in which the precipitation
5 chemicals are recovered from the sludge and recycled to the wastewater treatment.

In the treatment of wastewater, as a rule first mechanical separation of solid impurities is carried out, for instance with the aid of screens and grit chambers
10 and by allowing the solid impurities to settle in a preliminary settling device. Moreover, the wastewater is treated by chemical purification and preferably also by biological purification. The chemical purification is done such that precipitation chemicals, such as iron
15 salts or aluminium salts, are added to the water and, by flocculation, precipitate and collect impurities in the wastewater, such as phosphates and particles. In the biological purification, which can take place, for instance, by the activated sludge process or by means of
20 a trickling filter, the wastewater is purified by means of microorganisms. In wastewater treatment, large quantities of sludge are obtained, which must be taken care of. This can be carried out by digesting the sludge, in which case organic substances are converted into inorganic,
25 assisted by anaerobic microorganisms. The sludge obtained after digestion, i.e. digested sludge, can be used for landfilling or as fertiliser. If the digested sludge is to be used as fertiliser, its content of heavy metals, i.e. metals from the group consisting of chromium,
30 nickel, copper, zinc, cadmium, lead and mercury, should first be removed. Besides, the sludge contains the added precipitation chemicals, and from the economic point of view these should, if possible, be recovered and reused. In the present invention, precipitation chemicals relate

to iron and/or aluminium-containing compounds, such as ferric chloride, ferrous sulphate, ferric sulphate, aluminium sulphate and polyaluminium chloride.

Different methods for treating sludge from wastewater treatment are known, and as an example of prior-art technique, reference is made to WO96/20894, which was published on 11 July 1996. According to this reference, wastewater sludge is treated by acidifying the sludge to dissolve metals and phosphorus from the sludge. After separation of the remaining sludge, the precipitation chemicals iron and aluminium are recovered as phosphates by adjusting the pH to about 2-4. After separation of precipitated phosphates, a further precipitation is carried out, this time of dissolved heavy metals, which are precipitated by increasing the pH to about 7-9 and adding precipitants, such as sulphides. After separation, the heavy metal sulphides are deposited, while the filtrated water can be recycled to the wastewater treatment procedure. The resulting phosphate deposit, which contains iron phosphate and possibly also aluminium phosphates, can be treated for recovery of the precipitation chemicals iron and aluminium by adding an alkali hydroxide, such as sodium hydroxide, thereby forming insoluble iron hydroxide and a solution containing soluble alkali phosphates and aluminium hydroxide. The iron hydroxide can be dissolved in an acid, such as hydrochloric acid, sulphuric acid or nitric acid, to give a solution of the corresponding iron salt which is usable as precipitation chemical.

According to the above-mentioned WO 96/20894, the iron content of the sludge is present in trivalent form, or the iron is oxidised to trivalent form by adding an oxidant, such as hydrogen peroxide. No external addition of trivalent iron takes place. However, it is stated that external addition of phosphorus in the form of phosphoric acid or phosphate can take place to adjust the molar ratio of phosphoric acid to phosphorus to about 1:1.

As is apparent from that stated above, WO 96/20894 accomplishes a treatment of wastewater sludge, the sludge being relieved of undesired metals, such as heavy metals, and phosphorus. The content of metals originating from precipitation chemicals, such as iron and aluminium, in the sludge is recovered as phosphates and cannot be recycled directly to the wastewater purification process to be used as precipitation chemicals, but must first be converted by additional dissolving and precipitating procedures. Since each dissolving and precipitating procedure means a risk of decreased yield of the chemical at issue, it would be advantageous if a process could be provided, in which the metals used in the precipitation chemicals, after being dissolved from the wastewater sludge, can be recycled directly to the wastewater purification process, without any intermediate precipitating and dissolving steps.

According to the present invention, the above-mentioned drawbacks are obviated or reduced, and a method is provided for treating sludge from wastewater purification, in which iron and/or aluminium from the precipitation chemicals is dissolved from sludge, and the resulting solution is recycled to the wastewater treatment.

More specifically, the invention provides a method for treating sludge from wastewater purification, said sludge containing phosphorus and at least one metal which originates from precipitation chemicals and which is selected among Fe^{2+} and Al^{3+} ,

the pH of the sludge being adjusted to below 4 for dissolving the content of phosphorus and said metal in the sludge;

separating the remaining sludge;

treating the solution which is relieved of sludge and which contains phosphorus and said metal, for precipitation of the content of phosphorus in the solution as FePO_4 at a pH of 2-3; and

separating the precipitated FePO_4 . The invention is characterised in that the remaining solution, which contains said metal from the precipitation chemicals, is recycled to the wastewater treatment.

5 Further advantages and distinctive features of the invention will be evident from the following description and the appended claims.

The invention will now be described in more detail with reference to the accompanying drawing, which schematically shows a presently preferred embodiment of the
10 invention.

Sludge from a wastewater purification plant (not shown) containing, inter alia, phosphorus in the form phosphate and metals originating from the precipitation
15 chemicals that are used in the wastewater purification, is supplied to a first step I for dissolving the content of phosphorus and metals in the sludge from the precipitation chemicals. According to the invention, the metal or metals originating from the precipitation chemicals
20 are iron and/or aluminium, on the condition that iron is present in divalent form (Fe^{2+}). Originally, the iron is present in trivalent form (Fe^{3+}) in the precipitation chemical, but when the precipitation chemical in the chemical purification step of the wastewater treatment has
25 been added, flocculated and passed to the sludge phase, the iron is reduced to divalent form, for instance when digesting the sludge.

In the first step I, the content of phosphorus, iron and/or aluminium in the sludge is dissolved by acidifying
30 the sludge. This is effected by subjected the sludge to an acid hydrolysis at a pH below 4, preferably below 2, with an acid, e.g. sulphuric acid. The hydrolysis is effected under conditions that result in the desired dissolution. Neither temperature nor pressure is critical
35 in the hydrolysis, and ambient temperature and pressure can be used. If desired, an increased temperature and/or pressure, however, can be used in order to, for instance,

accelerate the hydrolysis. Normally the temperature can be in the range of about 0-200°C, and preferably the temperature is increased such as about 100-140°C, to accelerate the hydrolysis. Correspondingly, the pressure may vary from ambient pressure (atmospheric pressure) up to about 1 MPa depending on the hydrolysis temperature. It is in many cases sufficient for the pH to be just below 4 in the hydrolysis, but preferably the pH in the hydrolysis is below 2 for complete dissolution of the content of phosphorus, iron and/or aluminium in the sludge.

After completion of the hydrolysis, the remaining sludge and hydrolytic fluid are supplied to a second step II for separating the remaining sludge, for instance by filtration or centrifugation.

After separating the sludge, the solution relieved of sludge which contains dissolved phosphorus and metal from the sludge in the form of phosphate and dissolved metal salts, is supplied to a third step III for separating heavy metals, if any. By heavy metals are meant as mentioned above, metals from the group consisting of chromium, nickel, copper, zinc, cadmium, lead and mercury. If there are no heavy metals or if they can be neglected, this step can be omitted.

In the heavy metal separation step III, heavy metals are separated by adding a substance which forms an insoluble compound with heavy metals. Preferably, this substance is a sulphide ion source, such as sodium sulphide, such that the heavy metals are precipitated as heavy metal sulphides (HMS).

Alternatively, the content of phosphorus in the solution can first be precipitated as FePO_4 according to steps IV and V described below before the heavy metals are precipitated by adding a sulphide ion source.

If heavy metal sulphides can be accepted in the sludge which is separated after the acid hydrolysis, a further alternative implies that a sulphide ion source is

added even before or in connection with the acid hydrolysis for binding any heavy metals present as sulphides. In this case, the subsequent, specific sulphide precipitation step III can be omitted.

5 When the solution has been relieved of heavy metals, it is supplied to a fourth step IV, which is a step in which the pH of the solution is adjusted to 2-3, preferably 2-2.8. The pH is adjusted by adding a suitable base, such as sodium hydroxide or magnesium oxide. The adjustment of the pH is effected as a preliminary step before
10 the subsequent precipitation of the content of phosphorus in the solution as FePO_4 , which is insoluble in the stated pH range.

15 If the pH of the solution has already been adjusted, in the first step above, in which the pH is adjusted to below 4, to be in the range of 2-3, no subsequent adjustment of the pH is necessary after separating the sludge.

 The solution from step IV, which contains phosphate (PO_4^{3-}), divalent iron (Fe^{2+}) and/or aluminium (Al^{3+}), is
20 then supplied to a fifth step V for precipitating the phosphorus content of solution as iron phosphate (FePO_4). This takes place by adding to the solution a source of trivalent iron (Fe^{3+}), e.g. ferric chloride. With a view to achieving complete precipitation of the phosphorus
25 content of the solution, the trivalent iron is preferably added in an at least equimolar amount, i.e. in such an amount that the molar ratio of the trivalent iron to the phosphorus content of the solution is at least about 1:1, such as about 1-1.5:1. As mentioned above, trivalent iron
30 phosphate is insoluble in the pH range of 2-3, preferably 2-2.8, and, in this range, is precipitated in very pure form. To achieve as complete precipitation as possible of the phosphorus content of the solution, further a certain sojourn time should pass between the adding of the source
35 of trivalent iron and the separation of the formed iron phosphate. Suitably, the sojourn time is from about 5 min to about 6 h, preferably from about 30 min to about 1 h.

The precipitated iron phosphate is then removed from the solution in a per se known manner, for instance by filtration or centrifugation.

Even if the method has been described above such
5 that the step IV with pH adjustment of the solution is carried out before step V with the addition of a source of trivalent iron (Fe^{3+}), it should be understood that the relative order of steps IV and V is optional in the present invention. Thus it is quite possible, and in many
10 cases also preferred first to add the source of trivalent iron and only then to adjust the pH to 2-3. In the latter case, the above-mentioned sojourn time is placed in connection with the pH adjustment.

After separation of the precipitated iron phosphate,
15 the remaining solution contains the content of metal originating from the precipitation chemicals and being Fe^{2+} and/or Al^{3+} , in the original sludge. This solution, which is relieved of sludge, heavy metals and phosphate, is recycled to the wastewater purification for renewed use
20 of the content of iron and aluminium in the solution as precipitation chemicals. To make the iron content of the solution function actively as a precipitation chemical, it need be transferred from divalent to trivalent form. This is preferably carried out by the wastewater purification comprising an aerobic biological purification
25 step, and the solution being added to this purification step, the divalent iron being oxidated to trivalent iron in the aerobic biological purification step. If the recycled solution contains divalent iron, it will thus
30 for this reason be added to the wastewater purification prior to or in the aerobic biological purification step. If the solution contains aluminium only, it can in principle be added at an optional point in the wastewater purification. It goes without saying that it is also
35 possible to oxidise the divalent iron content of the solution to trivalent iron in some other manner, for instance by adding hydrogen peroxide. In this case, the

solution can be added to the wastewater purification at an optional point.

The ferric phosphate (FePO_4) resulting from the above described precipitation can be used as fertiliser in agriculture. It is also possible to recover the trivalent iron from the ferric phosphate for renewed use as precipitation reagent by treating the precipitation of ferric phosphate with an alkali, such as sodium hydroxide, to form ferric hydroxide which is separated and treated with an acid, such as hydrochloric acid or sulphuric acid, to form the corresponding ferric salt, which can then be used as precipitation reagent.

It will be appreciated that the invention offers a simple and smooth method for recovering, from sludge, metal originating from precipitation chemicals and recirculation of this metal to the wastewater purification for renewed use. By the metal, in the method, being constantly kept in solution, and there being no separation of the metal in one or more precipitation steps, the loss of the metal is minimised in the inventive method. The possibility of recovering the trivalent iron which constitutes precipitation reagent for the phosphorus content of the sludge also makes the inventive method being highly economic.

In addition to the recycling of iron and aluminium to the wastewater purification for renewed use as precipitation chemicals, a sludge is achieved by the present invention, which is relieved of undesired impurities and which can be used as, for instance fertiliser. Any heavy metals included are recovered in the invention preferably as a separate precipitate which can be deposited or be further processed for recovering the heavy metals. Finally, also the phosphorus content of the original sludge is recovered, according to the invention, separately in the form of ferric phosphate, which as stated above can be treated for recovering the trivalent iron. In this recovery of the trivalent iron, the phosphate is obtained as

sodium phosphate (Na_3PO_4) which, for instance, can be used as raw material for the production of fertiliser in agriculture or as raw material in the detergent industry.

As is obvious from that stated above, the emission of noxious or undesired substances is eliminated or reduced to a minimum level according to the invention, and consequently the invention provides an extremely environmentally friendly method for treating sludge from wastewater purification.

In order to further illustrate the invention, some embodiments will be described below, which, however, are not intended to limit the scope of the invention.

Example 1

Sludge from wastewater purification in a pilot plant was subjected to acid hydrolysis at a pH of 1.6 and a temperature of about 140°C for about 1 h. The precipitation chemicals comprised both ferric chloride and ferric sulphate and polyaluminium chloride, and the sludge therefore contained both Fe^{2+} and Al^{3+} . After the hydrolysis, the remaining sludge was separated by centrifugation, and the solution relieved of sludge (clear phase) was used in the test which was carried out at a temperature of about 20°C . A trivalent iron salt, which is stated in more detail in Table 1, was added to the solution under agitation to avoid settling in such an amount that the molar ratio $\text{Fe}^{3+}:\text{PO}_4^{3-}$ was 1:1. Then NaOH was added to the solution under continued agitation in order to adjust the pH of the solution to 2.6. In the pH adjustment, iron phosphate (FePO_4) was precipitated from the solution and after agitation and precipitation for 1 h, the resulting iron phosphate precipitation was separated from the solution by filtration through a GF/A filter. The remaining solution was then analysed in respect of the content of Fe^{2+} and Al^{3+} . The results are stated in Table 1. In Table 1, Fe^{2+} in and Al^{3+} in designate the content of Fe^{2+} and Al^{3+} , respectively, of the original sludge. Fe^{2+} out and Al^{3+} out designate the Fe^{2+} and Al^{3+} content of the

final, recycled solution. P recycled, Fe^{2+} recycled and Al^{3+} recycled designate the percentage amount of P, Fe^{2+} and Al^{3+} , respectively, which is recycled to the waste-water purification.

Table 1

Fe ³⁺ -source	Fe ³⁺ /P in (mole/mole)	P recycled Fe ²⁺ in (mg/l)	Fe ²⁺ out (mg/l)	Fe ²⁺ recycled Al ³⁺ in (mg/l)	Al ³⁺ out (mg/l)	Al ³⁺ recycled (%)
JKL 1)	1:1	726	631	87	20	74
PIX-111 2)	1:1	726	654	90	19	70
PIX-115 3)	1:1	726	670	92	20	74

1) JKL = iron chloride sulphate with 11.6% by weight Fe³⁺ and max. 20% by weight Cl⁻ and 20% SO₄²⁻.

JKL can be obtained from Kemira Kemwater, Helsingborg, Sweden.

2) PIX-111 = iron chloride with 13.7% by weight Fe³⁺ and 26-28% by weight Cl⁻.

PIX-111 can be obtained from Kemira Kemwater, Helsingborg, Sweden.

3) PIX-115 = iron sulphate with 11.5% by weight Fe³⁺ and 32% by weight SO₄²⁻.

PIX-115 can be obtained from Kemira Kemwater, Helsingborg, Sweden.

As appears from Table 1, the invention allows recycling of about 90% of Fe^{2+} and about 75% of Al^{3+} from the precipitation chemicals in the sludge.

Example 2

5 Wastewater sludge from a commercial wastewater purification plant which used ferric chloride and iron sulphate as precipitation chemicals, was subjected, in two different tests (Tests 1 and 2) to acid hydrolysis at a pH of 1.8 and a temperature of about 140°C for about 1 h.
10 After the hydrolysis, the remaining sludge was separated by means of centrifugal decanter, and the solution relieved of sludge (clear phase) was used in the test which was carried out at a temperature of about 50-60°C. A trivalent iron salt was added to the solution in a mixing tank such that the molar ratio of Fe^{3+} to PO_4^{3-} was about 1:1. The added trivalent iron was an iron chloride product containing 13.7% by weight Fe^{3+} and 26-28% by weight Cl^- . This product can be obtained from Kemira, Kemwater, Sweden, under the designation PIX-111. The
15 sojourn time in the mixing tank was 30 min. In a subsequent mixing tank, NaOH was added for adjusting the pH to 2.1-2.8. The sojourn time in this mixing tank was 30 min. In the pH adjustment, iron phosphate (FePO_4) was precipitated, which was separated by the solution from the mixing tank being pumped to a centrifugal decanter. A
20 cationic polymer Zetag 89 supplied by Allied Colloid, Great Britain, was added to ensure good separation in the decanter. An analysis was made of the Fe^{2+} content (Fe^{2+} in) of the original sludge and of the Fe^{2+} (Fe^{2+} out) content of the final solution. The results from Tests 1 and
25 2 are stated in Table 2, and the values concern average values in the tests, which were carried out during 4 h (Test 1) and 6 h (Test 2).

30 Table 2 shows that at least 80% of the Fe^{2+} content of the sludge can be recycled (Fe^{2+} recycled) to the water purification procedure.

Table 2

	Fe ³⁺ -source	Fe ³⁺ /P (mole/mole)	Fe ²⁺ in (mg/l)	Fe ²⁺ out (mg/l)	Fe ²⁺ recycled (%)
Test 1	PIX-111	1.07:1	56	47	84
Test 2	PIX-111	1.30:1	39.5	32.6	83

CLAIMS

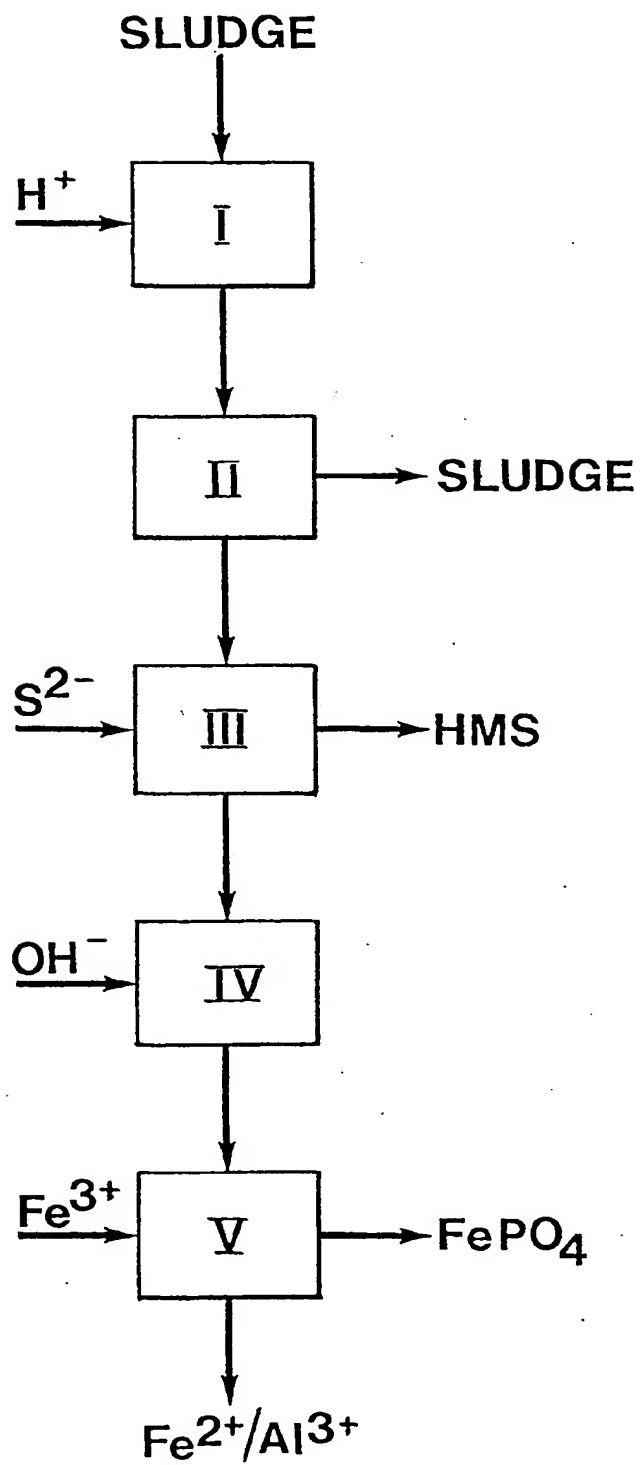
1. A method for treating sludge from wastewater
5 purification, said sludge containing phosphorus and at least one metal which originates from precipitation chemicals and which is selected among Fe^{2+} and Al^{3+} ,
the pH of the sludge being adjusted to below 4 for
dissolving the content of phosphorus and said metal in
10 the sludge;
separating the remaining sludge;
treating the solution which is relieved of sludge
and which contains phosphorus and said metal, for precipitation of the content of phosphorus in the solution as
15 FePO_4 at a pH of 2-3; and
separating the precipitated FePO_4 ;
characterised in that the remaining solution,
which contains said metal from the precipitation chemicals, is recycled to the wastewater treatment.
- 20 2. A method as claimed in claim 1, characterised in that the metal from the precipitation chemicals is Fe^{2+} .
3. A method as claimed in claim 1 or 2, characterised in that the pH of the sludge is adjusted
25 to below 2 for dissolving the content of phosphorus and said metal in the sludge.
4. A method as claimed in any one of claims 1-3, characterised in that the content of phosphorus in the solution is precipitated as FePO_4 by
30 adding at least the equivalent amount of Fe^{3+} .
5. A method as claimed in any one of claims 1-4, characterised in that the remaining solution contains Fe^{2+} and is recycled to wastewater treatment prior to an aerobic biological purification step.
- 35 6. A method as claimed in any one of claims 1-4, characterised in that the remaining solution

contains Fe^{2+} and is recycled to an aerobic biological purification step in the wastewater treatment.

7. A method as claimed in any one of claims 1-6, characterised in that heavy metals in the
5 sludge, which have been dissolved when adjusting the pH of the sludge to below 4 and which after separation of the remaining sludge is present in the solution relieved of sludge, are precipitated as sulphides by adding a sulphide ion source to the solution after separation of the
10 remaining sludge.

8. A method as claimed in claim 7, characterised in that the sulphide ion source is added to the solution before precipitation of the content of phosphorus in the solution.

15 9. A method as claimed in any one of claims 1-8, characterised in that the content of phosphorus and said metal which originates from the precipitation chemicals and which is present in the sludge is dissolved by acid hydrolysis at a temperature
20 in the range of 0-200°C.



1
INTERNATIONAL SEARCH REPORT

International application No. .

PCT/SE 98/00444

A. CLASSIFICATION OF SUBJECT MATTER

IPC6: C02F 11/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC6: C02F

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

WPI

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 9620894 A1 (KEMIRA CHEMICALS OY), 11 July 1996 (11.07.96), page 8, line 20 - line 31	1-3,5-9
Y	--	4
Y	EP 0530396 A1 (INDUSTRIE AUTOMATION SONDERTECHNIK GMBH & CO KOMMANDITGESELLSCHAFT), 10 March 1993 (10.03.93), claim 1	4
A	US 4416779 A (WILHELM K. RIPL ET AL), 22 November 1983 (22.11.83), claim 1	1-9
	--	

☒ Further documents are listed in the continuation of Box C.☒ See patent family annex.

* Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

29 June 1998

Date of mailing of the international search report

01-07-1998

Name and mailing address of the ISA/
Swedish Patent Office
Box 5055, S-102 42 STOCKHOLM
Facsimile No. +46 8 666 02 86

Authorized officer

Mårten Hulthén
Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 98/00444

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	Patent Abstracts of Japan, Vol 91, No 806 abstract of JP 31-14587 A (NISHIHARA KANKYO EISEI KENKYUSHO), 15 May 1991 (15.05.91) -- -----	1,3,4

INTERNATIONAL SEARCH REPORT
Information on patent family members

09/06/98

International application No.
PCT/SE 98/00444

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9620894 A1	11/07/96	AU 4306296 A	24/07/96
		CA 2208970 A	11/07/96
		CZ 9702032 A	12/11/97
		EP 0800483 A	15/10/97
		FI 97290 B,C	15/08/96
		FI 946190 D	00/00/00
		NO 972992 A	26/06/97
		PL 321541 A	08/12/97
EP 0530396 A1	10/03/93	DK 109492 A	05/03/93
		US 5296151 A	22/03/94
US 4416779 A	22/11/83	EP 0080981 A,B	08/06/83
		SE 441441 B,C	07/10/85
		SE 8205294 A	15/09/82